



Experimental Determination of the Impacts of Sediment Desiccation and Rewetting on Sediment Physical and Chemical Characteristics in Lawrence Lake, Pool 8, Upper Mississippi River

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PURPOSE: This technical note examines changes in sediment physical and chemical characteristics due to dessication and rewetting in laboratory systems. Results from this study may be used to better understand how sediment changes as a result of a lake drawdown and has implications for macrophyte growth potential in shallow lake rehabilitation programs.

BACKGROUND: Evidence is increasing that submersed and emergent macrophyte communities are a very important component of shallow lake ecosystems (Scheffer 1998 and references therein). Macrophyte communities minimize sediment resuspension by dampening wave activity (James and Barko 1994, Barko and James 1998); they also provide refugia for fish and zooplankton populations (Timms and Moss 1984; Killgore, Hoover, and Morgan 1991; Dibble, Killgore, and Harrel 1997) and substrate for invertebrate communities (Beckett, Aartila, and Miller 1992). Shallow lakes that have lost macrophyte communities usually exhibit frequent periods of sediment resuspension, high turbidity levels, low light penetration, enhanced nutrient recycling, and the prevalence of nuisance cyanobacteria (Bengtsson and Hellström 1992; James and Barko 1994; Scheffer et al. 1997; Bachmann, Hoyer, and Canfield 2000). While it is desirable to promote macrophyte growth to improve water quality in shallow systems, it is usually difficult to reestablish macrophyte communities without some form of manipulation to improve light conditions (i.e., reduce resuspension) and to remove destructive benthivorous fish such as carp (*Cyprinus carpio*). One potential manipulation to improve conditions for macrophyte growth in lakes that have lost this component is a lake drawdown to promote sediment consolidation and reduction in resuspension potential (Scheffer 1998). Fish stocks can also be manipulated during the drawdown to remove bioturbation impacts to the water column. When resuspension is controlled via this lake management technique, macrophyte communities can be successfully reestablished either by planting desirable species (Doyle and Smart 1993, Smart and Dick 1999) or by allowing the native seed bank in the sediments to germinate (Madsen et al., in preparation).

More information is needed regarding impacts of changes in sediment characteristics as a result of the desiccation and rewetting process on shallow lake ecosystem dynamics. Sediment desiccation and rewetting may result in positive impacts such as reduction in resuspension potential, improvement of rooting medium (i.e., nutrients and sediment texture) for macrophyte growth, conversion of soluble nutrients to particulate forms, and reductions in organic matter concentrations. However, negative impacts include conversion of organic matter to soluble N and P, enhanced nutrient release rates from sediments, and potential stimulation of algal growth. Ultimately, these positive and negative impacts of sediment desiccation and rewetting need to be evaluated with respect to the potential for improved conditions for macrophyte growth in shallow systems.

Pool 8, a run-of-the-river impoundment of the Upper Mississippi River system, is currently being considered for water level drawdown to consolidate sediments and stimulate submersed and

emergent macrophyte growth. Surficial sediments collected from a backwater region of this pool (Lawrence Lake) were experimentally dried and rewetted in the laboratory. Changes in sediment physical-chemical characteristics as a result of the desiccation-rewetting process and the implications of these changes to macrophyte growth in this system are reported herein.

METHODS: Lawrence Lake is a relatively large backwater region located on the Minnesota side of Pool 8, Upper Mississippi River (Figure 1). It currently has a diverse macrophyte species assemblage surrounding the perimeter of the lake. In June 2000, over 50 intact replicate sediment cores were collected at a station (depth= 0.7 m) located near the entrance to Lawrence Lake (Figure 1). The sediment samples were collected with a Wildco KB sediment core sampler (Wildlife Supply Co., Saginaw, MI) equipped with a plastic core liner (7.6 cm in diameter and 20 cm long). In the laboratory, the upper 8 cm of sediment were carefully extruded into a core liner that was precut to exactly 10 cm. A rubber stopper was used to seal the bottom of each extruded sediment core. The sediment core incubation systems were desiccated in the laboratory under controlled temperature conditions (~20 °C) in a darkened room. During the initial stages of the dessication process, only the surface of the sediment core was directly exposed to air. Drying and consolidation of the sediment core over time resulted in exposure of the sides of the core, as well as its surface, to air.

The experimental dessication and reflooding process is shown in Table 1. For each treatment, six sediment cores were sacrificed for determination of physical and chemical characteristics. In order to estimate the loss of moisture from sediment cores over time, the 10-cm core liner and rubber stopper were weighed before and immediately after extrusion of sediment into them. The loss of mass as a result of dessication was calculated by difference and converted to a percentage of the initial mass. For treatments 5 and 6 (Table 2), sediments were rewetted with 100 mLs of lake water (collected from Lawrence Lake) that had been previously filtered through a glass fiber filter (Gelman A/E). When filled with water, sediment cores were completely covered so that direct rehydration could occur via exposure of the top and sides of the sediment core. The sediment core incubation systems containing lake water were covered with plastic film to minimize evaporative losses. The volume of water remaining (i.e., not soaked up by the sediment) prior to sediment analysis was measured and analyzed for total and soluble reactive phosphorus (P), total nitrogen (N), ammonium-N ($\text{NH}_4\text{-N}$), and nitrate-nitrite-N ($\text{NO}_2\text{-NO}_3\text{-N}$) using standard analytical techniques (American Public Health Association (APHA) 1992; see below).

During the rewater phase, water surrounding the sediment cores was analyzed for dissolved oxygen, pH, and eH at 1- to 3-day intervals over a 2-week period. Dissolved oxygen (YSI BOD polarographic probe 5010 series) and pH were precalibrated against Winkler titrations (APHA 1992) and known pH buffer solutions. eH was measured using a combination platinum probe with an AgCl reference electrode (Fisher Acumet) referenced against saturated quinhydrone buffer solutions (pH 4 and pH 7 at 20 °C).

For each treatment, sediment core dimensions (height and diameter) were measured and the sediment was gently homogenized prior to analysis. Fresh sediment was carefully placed in a crucible and dried at 105 °C to a constant weight for determination of moisture content and sediment density (Allen et al. 1974), then combusted at 550 °C in a muffle furnace for determination of particulate

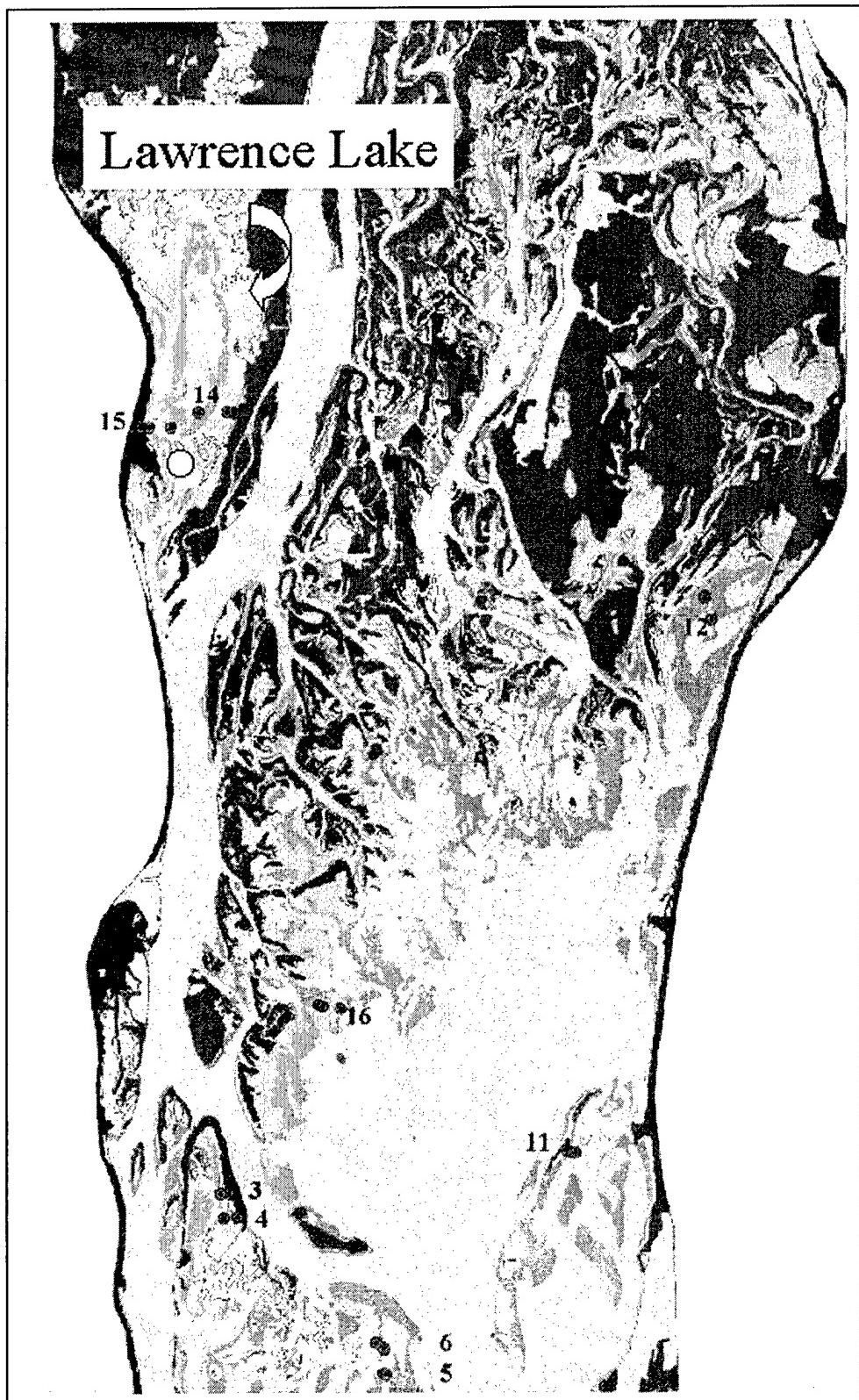


Figure 1. Sampling location in Lawrence Lake

Table 1
Mean¹ (1 S.E; n=6) Physical and Chemical Characteristics of
Sediment Collected in Lawrence Lake

Variable	Concentration
Moisture Content, %	80.0 (0.2)
Sediment Density, g/mL	0.212 (0.002)
Particulate Organic Matter, %	12.4 (0.2)
Porewater Soluble Reactive P, mg/L	2.506 (0.219)
Sediment Loosely Bound P, mg/g	0.022 (0.002)
Sediment Iron-bound P, mg/g	0.665 (0.024)
Sediment Aluminum-bound P, mg/g	0.210 (0.036)
Sediment Calcium-bound P, mg/g	0.064 (0.014)
Sediment Labile Organic P, mg/g	0.281 (0.031)
Sediment Refractory Organic P, mg/g	0.208 (0.032)
Sediment Total P, mg/g	1.450 (0.017)
Porewater Ammonium-N, mg/L	1.536 (0.355)
Porewater Nitrate-Nitrite-N, mg/L	0
Sediment Exchangeable Ammonium-N, mg/g	0.072 (0.004)
Sediment Organic N, mg/g	
Sediment Total N, mg/g	6.518 (0.091)
P Release Under Oxic Conditions, mg m ⁻² d ⁻¹	0.4 (0.1)
P Release Under Anoxic Conditions, mg m ⁻² d ⁻¹	9.2 (1.2)
NH ₄ -N Release Under Oxic Conditions, mg m ⁻² d ⁻¹	48.2 (8.4)
NH ₄ -N Release Under Anoxic Conditions, mg m ⁻² d ⁻¹	71.2 (3.4)
NO ₂ NO ₃ -N Release Under Oxic Conditions, mg m ⁻² d ⁻¹	11.2 (7.5)

¹ 1 S.E., n = 6.

Table 2
Description of Sediment Desiccation Timetable

Treatment #	Description
1	Initial conditions before the start of desiccation
2	50% dewatered (i.e., 32 days of dessication)
3	80% dewatered (i.e., 54 days of dessication)
4	100% dewatered (i.e., 70 days of dessication)
5	Rewetted for 1 day
6	Rewetted for 3 weeks

organic matter content (POM). Care was taken to minimize compaction of sediment in the crucible for density determination. Additional sediment was centrifuged at 4 °C at 3000 rpm for approximately 2 hr for separation of sediment porewater. The porewater was carefully decanted under a nitrogen atmosphere, filtered through a 0.45-µm filter (Gelman GN-6 Metrcel), and analyzed for soluble reactive P, NH₄-N, and NO₂NO₃-N using methods described in APHA (1992). Total sediment N and P concentrations were analyzed colorimetrically using Lachat QuikChem procedures (Lachat Method 10-107-06-2-D for N and 13-115-06-1-B for P; Zellweger Analytics, Lachat Div., Milwaukee, WI) following digestion with sulfuric acid, potassium sulfate, and red mercuric oxide (Plumb 1981). Sequential fractionation of inorganic P in the sediments was conducted according to Hielties and Lijklema (1980) and Nürnberg (1988) for the determination of ammonium-chloride-extractable P (NH₄Cl-P; loosely bound and CaCO₃-adsorbed P), bicarbonate-dithionite-extractable P (BD-P; iron-bound P), sodium hydroxide-extractable P (NaOH_{reactive}-P; aluminum-bound P), and hydrochloric acid-extractable P (HCl-P; calcium-bound P). A subsample of the NaOH extract was digested with potassium persulfate to determine unreactive NaOH-extractable P (NaOH_{nonreactive}-P; Psenner and Puckso (1988)). Labile organic P was calculated as the difference between NaOH_{reactive}-P and NaOH_{nonreactive}-P. Refractory organic P was calculated as total P minus labile organic P and inorganic P fractions. Each extraction was filtered through a 0.45-µm filter, adjusted to pH 7, and analyzed for soluble reactive P. The loosely bound P, iron-bound P, and labile organic P fractions in the sediments are classified as labile (i.e., exchangeable) sediment P, and the aluminum-bound P, calcium-bound P, and refractory organic P fractions as refractory (i.e., not exchangeable) sediment P (Penn et al. 1995). Exchangeable NH₄-N plus porewater NH₄-N were determined by cation exchange according to Bremner (1965). Exchangeable NH₄-N was corrected for porewater NH₄-N concentrations by difference. Sediment organic N was calculated as the difference between total N and the sum of exchangeable NH₄-N, porewater NH₄-N, and porewater NO₂NO₃-N.

N and P release from sediment under oxic and anoxic conditions was determined using additional sediment cores that were subjected to treatments 1 and 6. Experimental cores were transferred to a core liner (6.5-cm ID and 25-cm length) and 300 ml of filtered (Gelman A/E glass fiber) lake water were siphoned onto the sediment. The systems were sealed with rubber stoppers. For treatment 6,

consolidation of sediments resulted in a decrease in the sediment surface area and core length. Thus, it was much smaller than the diameter of the sediment incubation system. To prevent exposure of the sides of the core to overlying water during nutrient release analysis, the sides of the core were first gently wrapped in two layers of cellophane (leaving the top open), and then fine-grained sand was gently packed around the sides of the core after placement in the incubation system.

The redox environment in each system was controlled by bubbling the water with air (for oxic conditions) or nitrogen (for anoxic conditions). Sediment systems were incubated at 20 °C over a 1-week period to simulate in situ temperatures during summer. Samples were collected in the middle of the water column in each system on a daily basis for SRP and NH₄-N and NO₂NO₃-N analysis. Rates of N and P release from the sediment ($\text{mg m}^{-2} \text{ d}^{-1}$) were calculated as the linear change in concentration in the overlying water divided by time and the area of the incubation core liner. Changes in the surface area of the sediment core due to desiccation were factored into the rate calculations.

RESULTS: The surface sediments at the sampling station in Lawrence Lake exhibited a high moisture content and low sediment density, indicative of flocculent, fine-grained particles (Table 1). Sediment inorganic P fractions (i.e., loosely bound P, iron-bound P, aluminum-bound P, and calcium-bound P) accounted for 66 percent of the sediment total P, while sediment organic P fractions (i.e., labile and refractory organic P) accounted for 34 percent of the sediment total P. Iron-bound P was the dominant sediment inorganic P fraction (69 percent), followed by aluminum-bound P (22 percent), and calcium-bound P (7 percent). Loosely bound P accounted for only 2 percent of the sediment inorganic P fraction. Sediment total N was dominated by the organic N fraction (98 percent; Table 1). Sediment exchangeable NH₄-N accounted for approximately 1 percent of the sediment total N. Porewater concentrations of NH₄-N and SRP, as well as rates of NH₄-N and P release under anoxic conditions, were high relative to other eutrophic aquatic systems (Barko and Smart 1986; Nurnberg et al. 1986).

During the dessication process (i.e., treatments 1-4), sediment cores lost water at a constant rate of 1.5 percent water mass d^{-1} (Figure 2). Complete dryness occurred in approximately 70 days. Sediment cores consolidated as a consequence of water loss, resulting in a nearly constant decline in the sediment core length from approximately 8.5 cm to less than 4 cm (Figure 2). Core diameter declined from approximately 6.5 cm for treatment 1 to 4.3 cm for treatment 6.

During the rewetting process, the lake water used to rehydrate the sediment cores exhibited detectable dissolved oxygen during the first 2 days of exposure (Figure 3). Dissolved oxygen declined to less than 0.5 mg/L after day 2 of exposure. The eH of the lake water was greater than 75 mV during the first 2 days of exposure to desiccated sediment. In conjunction with low dissolved oxygen, it declined to less than -100 mV after day 2 of the rewetting process. The pH of lake water remained nearly constant at about 7.1 throughout the rewetting procedure.

The actual mean moisture content (i.e., the percentage of moisture as a function of sediment fresh weight as opposed to the percentage of water lost from the cores) declined, while mean sediment density increased nearly 7 times during the dessication process (Figure 4). Moisture content increased, while sediment density declined, as a result of rewetting the sediment cores for 1 day and

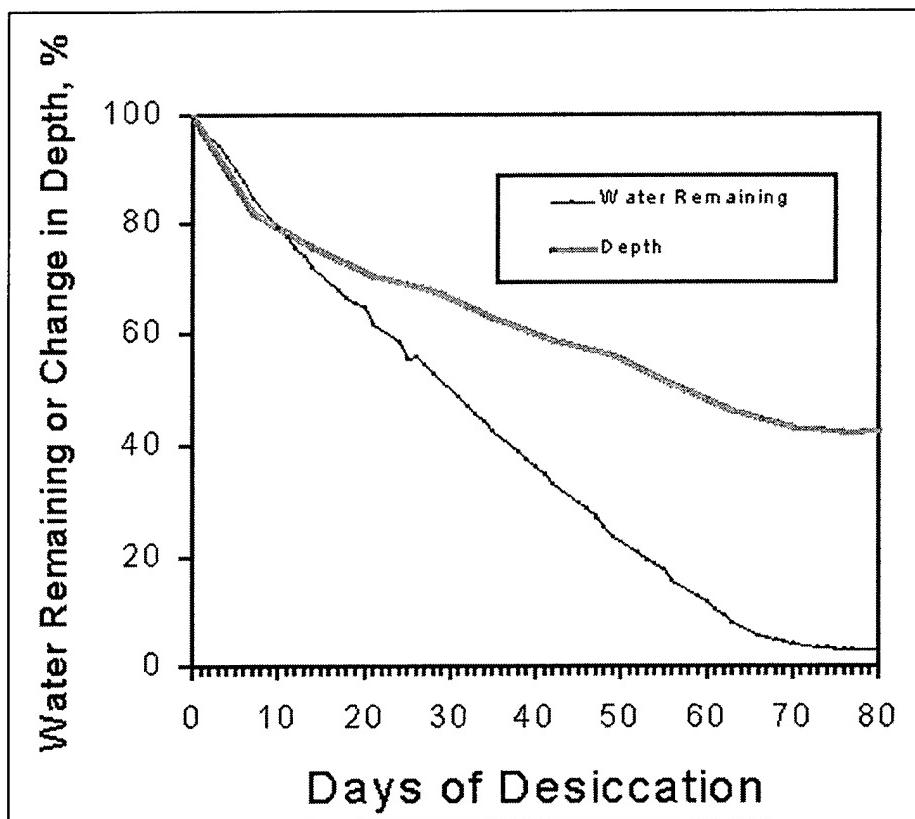


Figure 2. Changes in the percent water remaining and percent change in sediment core depth as a result of experimental desiccation of sediment cores measuring 8 cm long by 8 cm diameter

for 3 weeks (i.e., treatments 5 and 6). However, the mean moisture content of the sediment cores after the dessication-rewetting process was significantly lower (i.e., less than 40 percent; $p < 0.05$), compared to the initial mean moisture content of 80 percent (Table 1 and Figure 4). Similarly, mean sediment density was significantly higher ($p < 0.05$) after the dessication-rewetting process, compared to its initial mean concentration (Figure 4). Mean particulate organic matter content declined slightly ($p < 0.05$) from its initial mean concentration as a result of the dessication-rewetting process.

Sediment dessication and rewetting resulted in marked changes in sediment P characteristics (Figure 5). The mean mass of porewater P declined dramatically during the dessication process. It increased slightly as a result of rewetting the sediment cores; but there was an overall net loss of porewater P mass as a result of the dessication-rewetting process. The mean mass of loosely bound P increased substantially during the dessication process, then declined slightly as a result of rewetting the sediment cores. Overall, however, there was a net increase in loosely bound P as a result of the dessication-rewetting process (i.e., treatment 1 versus treatment 6). The mean mass of aluminum-bound P and calcium-bound P remained nearly constant during the dessication process. However, rewetting of the sediment cores coincided with a significant increase in the mean mass of these constituents. In contrast, the mean mass of iron-bound P remained constant during the dessication

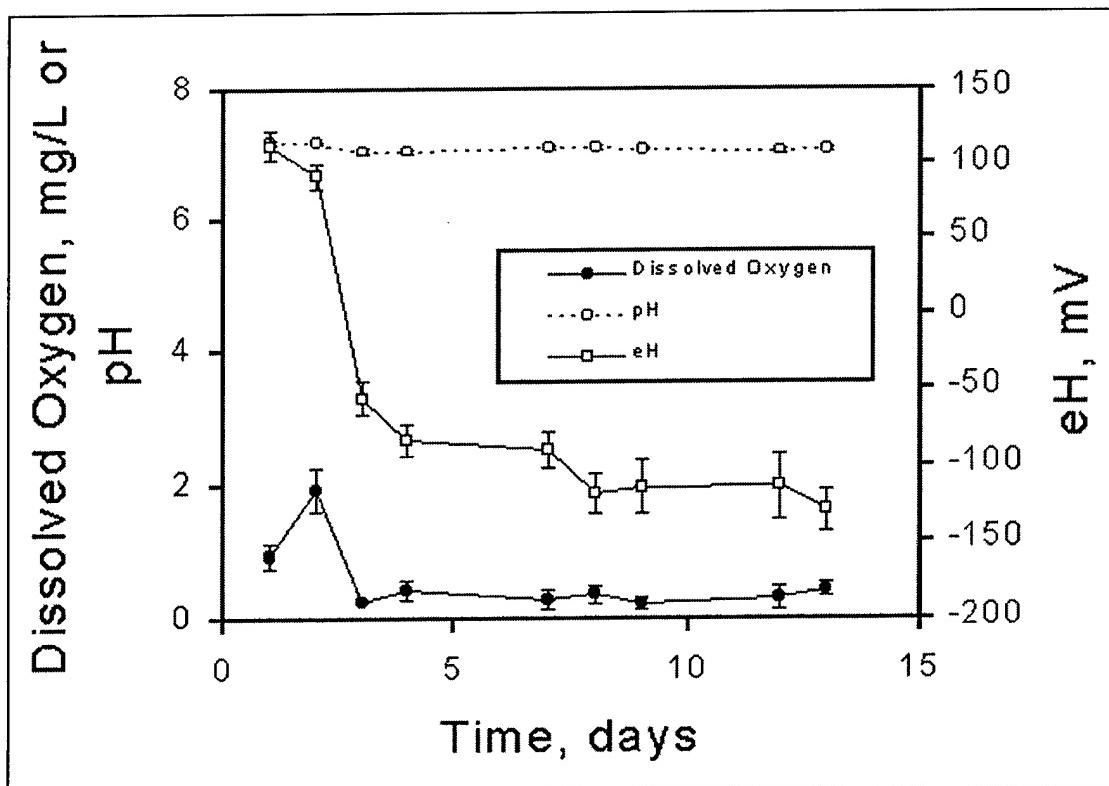


Figure 3. Variations in dissolved oxygen, pH, and eH of filtered lake water used to rehydrate desiccated sediment cores

process and declined significantly ($p < 0.05$) as a result of the rewetting process. The mean mass of sediment organic P (both labile and refractory organic P fractions) appeared to remain approximately constant throughout the dessication-rewetting process.

Overall net changes in sediment P fractions (i.e., treatment 1 versus treatment 6) as a result of the dessication-rewetting process are shown in Table 3. In general, sediment total P and organic P mass did not change significantly ($p > 0.05$) as a result of the dessication-rewetting process (Table 3). In contrast, loosely bound P, aluminum-bound P, and calcium-bound P exhibited a net increase in mass ($p < 0.05$) while iron-bound P exhibited a nearly equivalent net decrease in mass ($p < 0.05$). Less than 0.25 mg total P was measured in the excess water that was not soaked up by the sediment during the rewetting process, suggesting that loss of P mass via diffusion and desorption out of the core was minor in relation to exchanges of P between the various sediment P fractions.

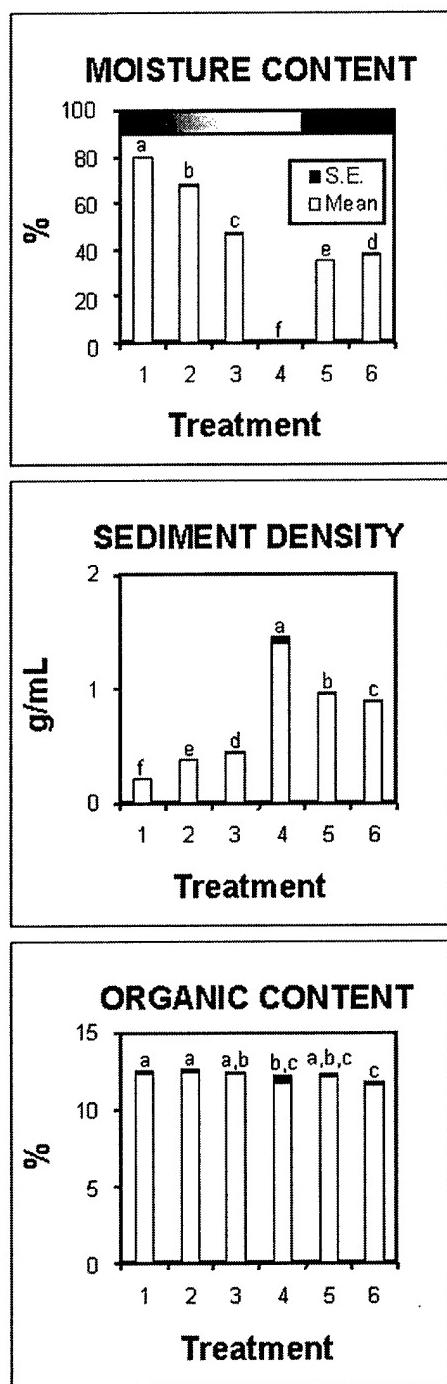


Figure 4. Variations in mean and standard error (SE; n=6) sediment moisture content, sediment density, and organic matter content of the sediment as a function of treatment. Treatment 1 = initial conditions, 2 = 50 percent dewatered, 3 = 80 percent dewatered, 4 = 100 percent dewatered, 5 = rewetted for 1 day, and 6 = rewetted for 3 weeks. Shaded bar indicates relative dryness of sediment cores (black = wet; white = dry). Different letters indicate significant differences in means ($p < 0.05$) based on Duncan's Multiple Range Analysis (ANOVA; SAS 1994)

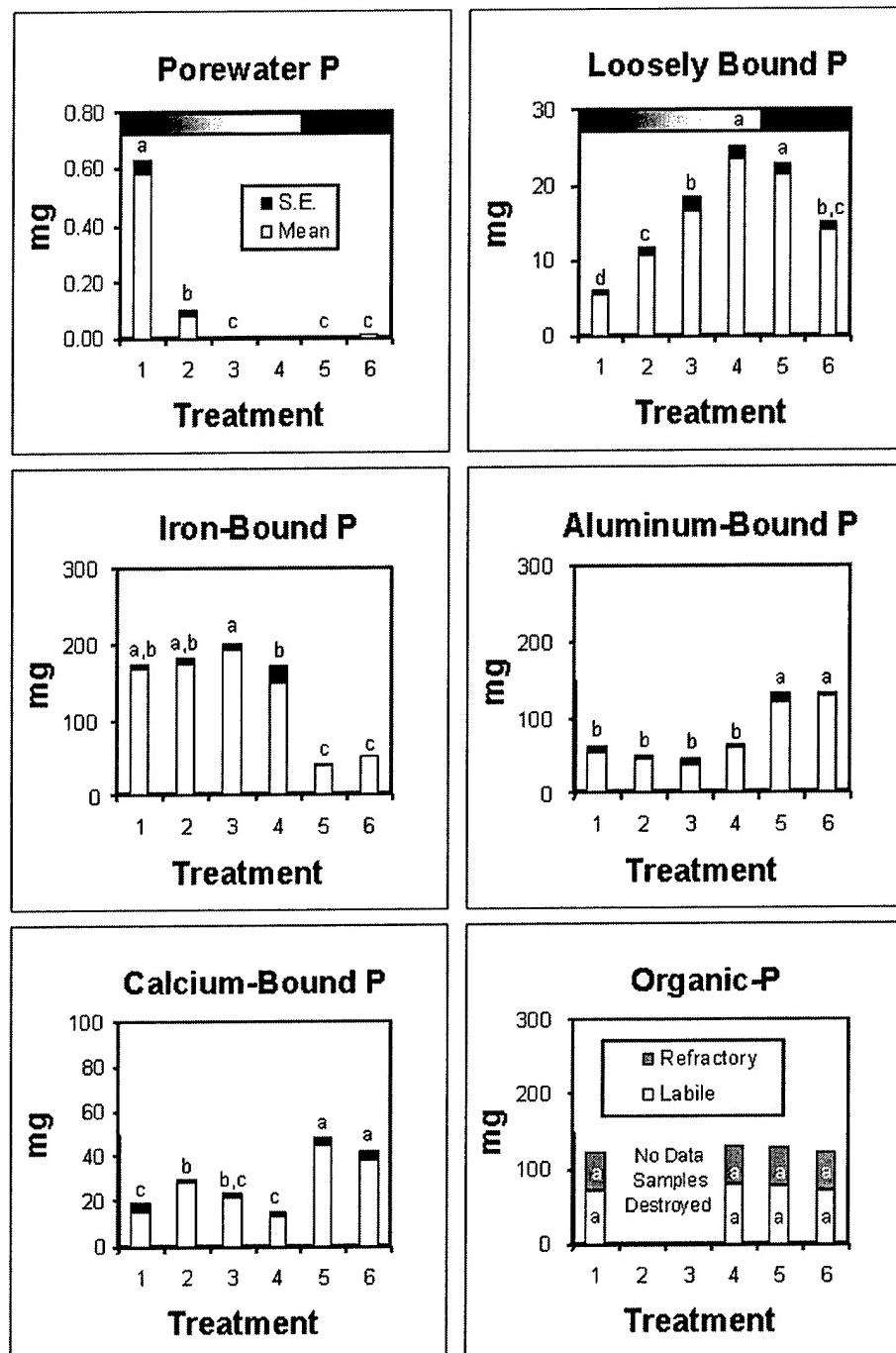


Figure 5. Variations in mean and standard error (SE; n=6) porewater phosphorus (P), Loosely bound P, iron-bound P, aluminum-bound P, calcium-bound P and organic P of the sediment as a function of treatment. Treatment 1 = initial conditions, 2 = 50 percent dewatered, 3 = 80 percent dewatered, 4 = 100 percent dewatered, 5 = rewetted for 1 day, and 6 = rewetted for 3 weeks. Shaded bar indicates relative dryness of sediment cores (black = wet; white = dry). Different letters indicate significant differences in means ($p < 0.05$) based on Duncan's Multiple Range Analysis (ANOVA; SAS 1994).

Table 3
Changes in the Mean¹ (1 S.E.) Mass Sediment P Constituents as a Result of the Desiccation-Rewetting Process²

Variable	Net Change (mg)
Porewater P	-0.6 (± 0.1)*
Loosely bound P	8.8 (± 1.0)*
Iron-bound P	-117.1 (± 6.4)*
Aluminum-bound P	75.7 (± 13.0)*
Calcium-bound P	22.2 (± 7.1)*
Labile Organic P	0.7 (± 9.5)
Refractory Organic P	-1.9 (± 15.1)
Total Sediment P	-11.8 (± 10.0)

¹ Treatment 1 versus treatment 6.

² Asterisks indicate significant changes ($p < 0.05$) mass (t-test, SAS (1994)).

Changes in sediment N fractions during the dessication and rewetting process are shown in Figure 6. The mean mass of porewater NH₄-N and sediment exchangeable NH₄-N declined significantly ($p < 0.05$) during dessication. However, the mean mass of these constituents increased substantially as a result of rewetting (Figure 6). Although there was no detectable porewater NO₂NO₃-N during the dessication process, NO₂NO₃-N in the porewater exhibited a slight increase as a result of the rewetting process. The mean mass of sediment organic N generally declined throughout the dessication/rewetting process.

Overall, mean sediment total N exhibited a statistically significant net loss of mass as a result of the dessication-rewetting process, which represented 19 percent of the original mean sediment N mass (Table 4). Total N mass measured in the excess water removed from the cores after the rewetting process (i.e., 1.3 mg) did not account for this net loss in sediment total N. Coincident with this pattern was a net decline in the mean mass of sediment organic N. There was no net change in the mean mass of porewater NO₂NO₃-N over the dessication-rewetting process. Although porewater NH₄-N and sediment exchangeable NH₄-N exhibited a net increase in mean mass as a result of the dessication-rewetting process, these constituents did not account for losses observed in sediment total and organic N.

As an apparent result of the dessication-rewetting process, mean rates of P release from sediments under anoxic conditions, measured in laboratory incubation systems, declined significantly ($p < 0.05$) over mean rates measured before treatment (Figure 7). This pattern coincided with declines in porewater concentrations of P (Figure 5). Mean rates of P release from sediments were greater under anoxic conditions than oxic conditions both before and after the dessication-rewetting process. The mean rate of P release from sediments under oxic conditions did not change as a result of the

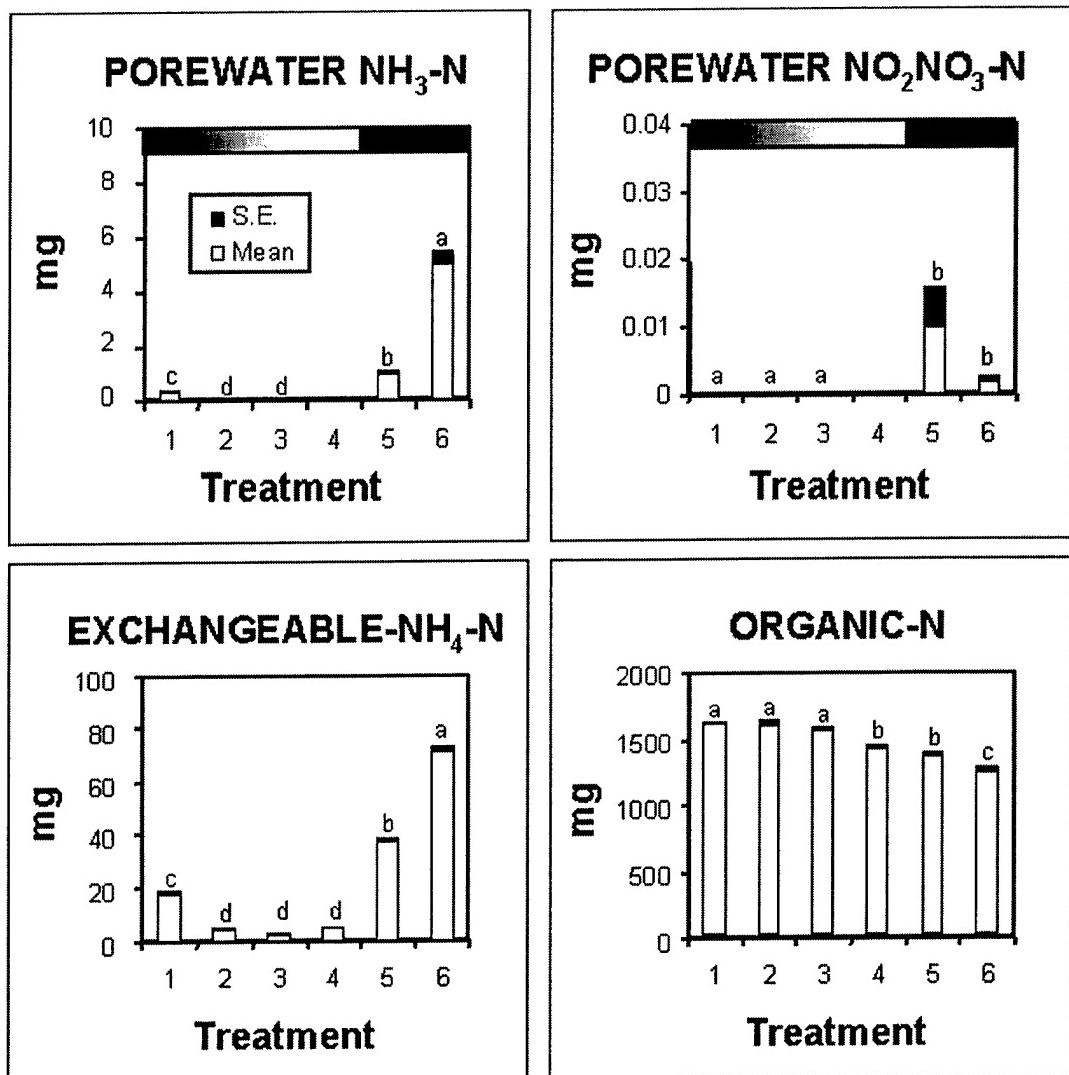


Figure 4. Variations in mean and standard error (SE; n=6) porewater ammonium (NH₄-N), porewater nitrate-nitrite (NO₂NO₃-N), exchangeable NH₄-N, and organic nitrogen (N) of the sediment as a function of treatment. Treatment 1 = initial conditions, 2 = 50 percent dewatered, 3 = 80 percent dewatered, 4 = 100 percent dewatered, 5 = rewetted for 1 day, and 6 = rewetted for 3 weeks. Shaded bar indicates relative dryness of sediment cores (black = wet; white = dry). Different letters indicate significant differences in means ($p < 0.05$) based on Duncan's Multiple Range Analysis (ANOVA; SAS 1994)

dessication-rewetting process. Mean rates of NH₄-N release from sediment increased substantially under both oxic and anoxic conditions after the dessication-rewetting process, compared to mean rates of N release from sediments measured before treatment (Figure 7). Mean rates of NO₂NO₃-N release under oxic conditions also increased substantially after the dessication-rewetting process (Figure 7). These patterns coincided with increases in concentrations of porewater and exchangeable NH₄-N and porewater NO₂NO₃-N after rewetting (Figure 6).

Table 4
Changes in the Mean (1 S.E.) Mass¹ Sediment N Constituents as a Result of the Desiccation-Rewetting Process²

Variable	Net Change (mg)
Porewater NH ₄ -N	4.6 (± 0.4)*
Porewater NO ₂ NO ₃ -N	< 1
Exchangeable NH ₄ -N	58.7 (± 1.4)*
Organic N	-379.3 (± 23.5)*
Total Sediment N	-316.0 (± 24.4)*

¹ Treatment 1 versus Treatment 6.

² Asterisks indicate significant changes ($p < 0.05$) mass (t-test, SAS (1994)).

DISCUSSION: The experimental desiccation process resulted in substantial sediment consolidation, as the percent moisture content and organic matter content declined, while sediment density increased after the rewetting process. Similarly, James et al. (2001) found that moisture and organic matter content decreased, while sediment density increased, following drawdown of Big Muskego Lake for purposes of sediment desiccation and consolidation. These changes in sediment characteristics need to be considered within the context of efforts to stimulate macrophyte growth in shallow aquatic systems. In particular, Barko and Smart (1986) demonstrated experimentally that low sediment density and high organic matter content suppressed macrophyte growth, while higher density sediment with lower organic matter content were associated with greater macrophyte growth response. Higher density sediments may provide a better plant rooting medium for macrophytes, resulting in increased root contact with sediment particles and associated nutrients (Barko and Smart 1986).

Sediment consolidation via desiccation may also indirectly enhance macrophyte growth potential by increasing the critical shear stress required to resuspend sediments (Scheffer 1998). Frequent resuspension of flocculent sediments and resulting high turbidity in shallow, wind-exposed lakes often leads to light-limiting conditions for macrophyte growth (Blom, Van Diun, and Vermaat 1994). For instance, using macrophyte growth models, Best, Boyd, and James (2001) suggested that the growth of *Vallisneria americana* and *Potamogeton pectinatus* becomes limiting at turbidity values in excess of 15-18 NTU in shallow Peoria Lake, IL. Because resuspension is frequent in this lake due to a low critical shear stress of sediments in relation to wind-generated shear stress (James, Best, and Barko, in preparation), turbidity usually exceeds the threshold required for macrophyte growth in this system. Thus, the reproductive survival and persistence of submersed species can decline substantially as a result of high turbidity (Doyle and Smart 2001; Kimber, Korschgen, and Van der Valk 1995), leading to population extinction. By increasing the critical shear stress required to resuspend sediments via desiccation and compaction to lessen the frequency of resuspension and periods of low light conditions, light conditions may improve substantially for successful macrophyte growth.

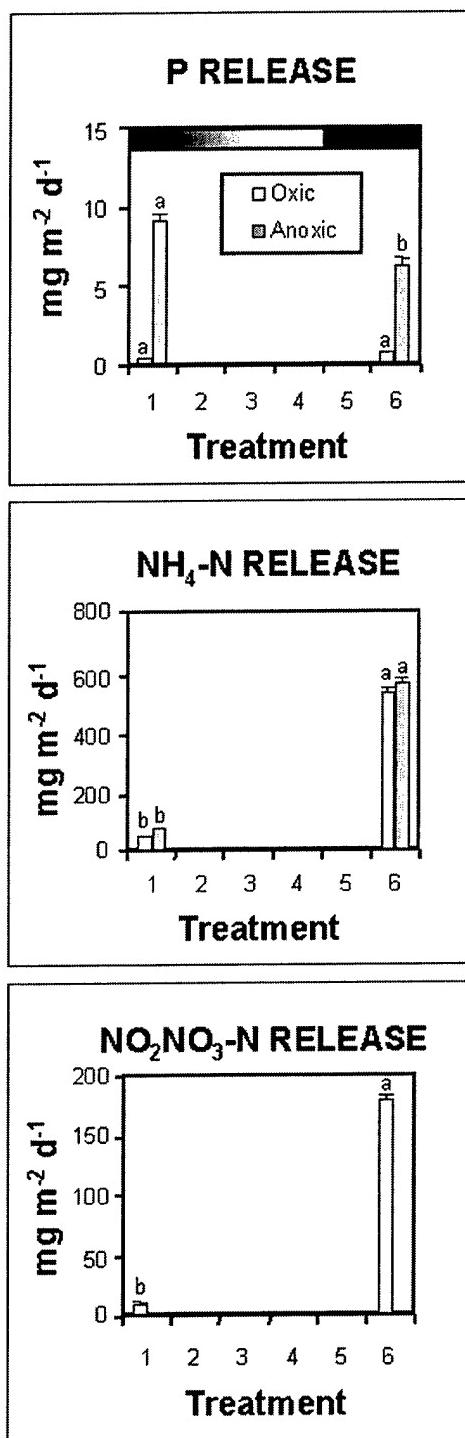


Figure 7. Variations in mean and standard error (SE; n=6) rates of phosphorus (P), ammonium-N ($\text{NH}_4\text{-N}$), and nitrate-nitrite-N ($\text{NO}_2\text{-NO}_3\text{-N}$) for treatments 1 and 6. Treatment 1 = initial conditions and 6 = rewetted for 3 weeks. Shaded bar indicates relative dryness of sediment cores (black = wet; white = dry). Different letters indicate significant differences in means ($p < 0.05$) based on Duncan's Multiple Range Analysis (ANOVA; SAS 1994).

Dynamic changes occurred in P associated with sediment as a result of sediment dessication and rewetting. It appeared that the loosely bound, aluminum-bound, and calcium-bound P fractions increased in conjunction with declines in the iron-bound P fraction after dessication and rewetting. In particular, 84 percent of the change in the iron-bound P fraction could be accounted for by increases in the more refractory aluminum- and calcium-bound P fractions, suggesting that dessication and rewetting of these sediments was associated with a net loss of sediment P from the labile P pool. Consistent with this change in the labile P pool of the sediments were declines in porewater concentrations of P and the rate of P release from the sediments under anoxic conditions. These patterns suggested that potential P flux from the sediments via diffusive processes was suppressed due to irreversible binding of P to aluminum and calcium (i.e., apatite formation) during the dessication-rewetting process.

In contrast, others have found that sediment desiccation resulted in an increase in iron-bound P. De Groot and Van Wijck (1993a, 1993b) attributed this pattern to oxidation of iron and sorption of P onto these particles during the desiccation process. They also suggested that some of the P mineralized from the organic P fraction was scavenged by iron and calcium in the sediment. In particular, they observed a decrease in both refractory and labile organic P fractions as a result of the desiccation process, suggesting enhanced microbial activity as a result of oxidation of sediments. Surprisingly, a statistically significant decline was not observed in either the labile or refractory sediment organic P fraction, nor was there an increase in the iron-bound P fraction as a result of desiccation and rewetting. The experimental drying process may have inhibited microbial activity and breakdown of organic P, compared to *in situ* desiccation of sediments in the De Groot and Van Wijck (1993a, 1993b) study.

Reasons for the marked increases in the aluminum and calcium-bound P fractions as a result of desiccation and rewetting of sediments from Lawrence Lake are not completely known. Perhaps the experimental drying and compaction process induced an early diagenesis of some of the sediment P (Penn et al. 1995), resulting in the conversion of some labile P into more refractory P (i.e., P that is not exchangeable and eventually buried; aluminum- and calcium-bound P). Whereas dessication of *in situ* lake and river sediments is usually subjected to temporary periods of rewetting due to precipitation, higher water levels, etc., sediments in this study were completely dried and compacted in an uninterrupted manner. This more complete desiccation pattern perhaps led to conversion of a portion of the labile pool to refractory P.

Changes in sediment P pools as a result of sediment desiccation appeared to provide both negative and positive impacts to macrophyte growth potential. Although P is generally not limiting to macrophyte growth, macrophytes derive most of their nutritional needs for this nutrient via root uptake from the sediments. Barko and Smart (1980) demonstrated that both porewater and labile P pools are accessed via root uptake. Since sediments from Lawrence Lake were rich in P, declines in the labile sediment P pool as a result of desiccation and rewetting would likely not impact macrophyte growth in this system, since concentrations are still high relative to growth needs. However, in other aquatic systems, declines in the available sediment P pool to limiting quantities could negatively impact macrophyte growth.

Positive impacts of a net decline in the labile sediment P pool and flux of P from the sediment after desiccation include a reduction in the flux of sediment-derived P to the water column for algal uptake and growth. Reduction in P flux from the sediment may impart a competitive advantage to submersed macrophytes for successful growth via increasing light penetration due to less algal biomass (and turbidity) in the water column. In contrast to patterns for sediments in Lawrence Lake, others have found that dessication can lead to substantial increases in the flux of P from sediments. In Big Muskego Lake, for instance, P flux from the sediment increased by one order of magnitude after desiccation and rewetting (James et al. 2001). Reasons for differences in the response of sediment P flux to the desiccation process for different sediments are not known and need to be investigated. In particular, changes in equilibrium processes (i.e., adsorption and desorption; De Groot and Van Wijck (1993b)) need to be examined in relation to changes in sediment P pools for different sediment types.

Sediment dessication and rewetting resulted in significant increases in sediment exchangeable NH₄-N, porewater NH₄-N and NO₂NO₃-N, and rates of NH₄-N and NO₂NO₃-N release from sediments. These changes are of ecological significance to macrophyte growth potential since root uptake is the principal means of N acquisition and macrophytes are usually limited by the availability of sediment N (Gerloff and Krombholz 1966; Barko et al. 1988; Barko, Gunnison, and Carpenter 1991; Rogers, McFarland, and Barko 1995). In particular, NH₄-N and exchangeable NH₄-N are both readily taken up via root systems. Anaerobic decomposition of organic N to NH₄-N was likely, because although dissolved oxygen in the water used to rewet the sediment was high (i.e., 1-2 mg/L) during the first day of rewetting, it subsequently declined to near zero as it soaked into the sediment, indicating that anoxic conditions were occurring during most of the rewetting process.

Initial rewetting (i.e., treatment 5) resulted in a small, but significant, increase in porewater NO₂NO₃-N, suggesting some oxidation of NH₄-N (i.e., nitrification) during the initial stages of the rewetting process. The initial introduction of oxygen via rewetting would provide a mechanism for conversion of NH₄-N to NO₂NO₃-N in the porewater via bacterial nitrification. The subsequent decline in NO₂NO₃-N and increase in NH₄-N in the porewater after 3 weeks of rewetting was likely the result of dissimilatory reduction of nitrate. Declines in porewater NO₂NO₃-N were attributed to denitrification, since introduced water was anoxic after the second day. eH declined to between zero and -125 mV during this time period, which was within the range for nitrate reduction (Tomaszek 1995).

When rewetted sediments were subjected to an oxic environment in laboratory incubation systems for analysis of N and P release from sediments (i.e., overlying water was added to the rewetted sediments and gently bubbled with air), rates of both NH₄-N and NO₂NO₃-N release from sediments increased substantially over initial rates observed prior to the desiccation process. The substantial increases in porewater NH₄-N and exchangeable NH₄-N observed in the sediment as a result of rewetting, and anaerobic metabolism of organic N, perhaps caused a shift in NH₄-N equilibrium toward flux out of the sediment and into the water column of the sediment incubation systems subjected to an oxic environment. Increases in the rate of NO₂NO₃-N release from rewetted sediments under an oxic environment were likely due to nitrification of some of the NH₄-N.

The most unusual feature of sediment N dynamics during the desiccation-rewetting process was an unexplained net loss of sediment N, primarily in the form of organic N, that could not be accounted for by conversion to exchangeable NH₄-N in the sediments or to NH₄-N or NO₂NO₃-N in the pore water. While more research is needed to identify and quantify the transformations that may be resulting in this net loss of N during desiccation-rewetting, possibilities include nitrification followed by denitrification (for NO₂NO₃-N; Mitchell and Baldwin (1999)) and/or volatilization of NH₄-N to NH₃. The pH of water used to rewet the sediment was near 7.0, suggesting that conversion to NH₃-N was minor. Downward shifts in the eH of water and the occurrence of anoxia during the rewetting process implicated the possibility of N loss via denitrification. However, the processes resulting in loss of sediment N need to be identified because if desiccation is leading to net N loss from the sediment, pool elevation drawdowns may have management implications for N regulation in the Upper Mississippi River system.

SUMMARY: Experimental desiccation and rewetting of sediments from Lawrence Lake in Pool 8 of the Upper Mississippi River resulted in a decline in moisture content and organic matter, and an increase in sediment density, indicating consolidation of sediment. Pore water and iron-bound P, and rates of P release under anoxic conditions declined, while aluminum- and calcium-bound P increased as a result of desiccation-rewetting, resulting in a net increase in the sediment refractory P pool. Labile and refractory organic P in the sediment did not change as a result of the desiccation-rewetting process. Pore water NH₄-N and NO₂NO₃-N and exchangeable NH₄-N increased substantially as a result of the desiccation-rewetting process. Coincident with these constituent increases after rewetting were markedly elevated rates of NH₄-N and NO₂NO₃-N release from the sediment. However, there was an overall net loss of organic N as a result of the desiccation-rewetting process that could not be accounted for by increases in other N fractions. This pattern suggested that N was being lost to the atmosphere via denitrification as a result of the desiccation-rewetting process. Increases in available N, coupled with consolidation of flocculent sediments, suggested that desiccation of sediment in Lawrence Lake will likely result in improved conditions for macrophyte growth.

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